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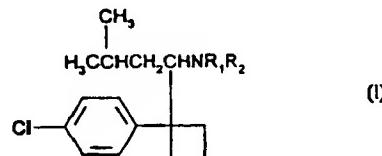
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(54) Title: METHOD OF TREATING SEXUAL DYSFUNCTION



(57) Abstract

A compound of formula (I), or a pharmaceutically acceptable salt thereof in which R₁ and R₂ are independently H or methyl (for example N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl amine hydrochloride optionally in the form of its monohydrate), including individual enantiomers and racemates thereof, is used for treating sexual dysfunction.

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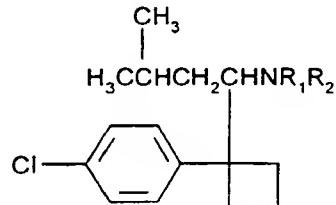
Method of Treating Sexual Dysfunction

This invention relates to a method of treating sexual dysfunction.

5

Sexual dysfunction includes, but is not limited to, the following: impotence, that is the inability to copulate and in a male may involve an inability to achieve penile erection or ejaculation or premature ejaculation. Various drugs have been suggested for the treatment of this condition, including paroxetine and 10 sildenafil (Viagra®). However, there exists a need for efficacious compounds which have a low profile of side effects.

According to the present invention there is provided a method of treating sexual dysfunction, in which a therapeutically effective amount of a compound of 15 formula I



including enantiomers and pharmaceutically acceptable salts thereof, in which R_1 , 20 and R_2 are independently H or methyl, is administered in conjunction with a pharmaceutically acceptable diluent or carrier to a human in need thereof.

The compounds of formula I are advantageous in that they may be used to provide a higher degree of efficacy and a lower degree of side effects 25 compared to the conventional treatments.

A preferred compound of formula I is N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine or a salt thereof, for example the hydrochloride salt. A preferred form of this hydrochloride is its monohydrate.

- 5 The preparation and use of compounds of formula I, such as N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine and salts thereof, in the treatment of depression is described in British Patent Specification 2098602. The use of compounds of formula I such as N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine and salts thereof in the treatment of 10 Parkinson's disease is described in published PCT application WO 88/06444. The use of N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine and salts thereof in the treatment of cerebral function disorders is described in US Patent 4939175. The use of N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride in the treatment of obesity is described in 15 published PCT application WO90/06110. A particularly preferred form of this compound is N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride monohydrate (sibutramine hydrochloride) which is described in European Patent Number 230742. The use of N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine and salts thereof for improving the 20 glucose tolerance of humans having Impaired Glucose Tolerance or Non-Insulin Dependent Diabetes Mellitus is described in published PCT application WO95/20949.

- 25 It will be appreciated by those skilled in the art that compounds of formula I contain a chiral centre. When a compound of formula I contains a single chiral centre it may exist in two enantiomeric forms. The present invention includes the use of the individual enantiomers and mixtures of the enantiomers. The enantiomers may be resolved by methods known to those skilled in the art, for example by formation of diastereoisomeric salts or complexes which may be 30 separated, for example, by crystallisation; via formation of diastereoisomeric derivatives which may be separated, for example, by crystallisation, gas-liquid or liquid chromatography; selective reaction of one enantiomer with an enantiomer-

specific reagent, for example enzymatic oxidation or reduction, followed by separation of the modified and unmodified enantiomers; or gas-liquid or liquid chromatography in a chiral environment, for example on a chiral support, for example silica with a bound chiral ligand or in the presence of a chiral solvent. It
5 will be appreciated that where the desired enantiomer is converted into another chemical entity by one of the separation procedures described above, a further step is required to liberate the desired enantiomeric form. Alternatively, specific enantiomers may be synthesised by asymmetric synthesis using optically active reagents, substrates, catalysts or solvents, or by converting one enantiomer to
10 the other by asymmetric transformation.

Preferred compounds of formula I are N,N-dimethyl-1-[1-(4-chlorophenyl)-cyclobutyl]-3-methylbutylamine, N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N- methylamine, and 1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine including racemates, individual enantiomers and mixtures thereof, and pharmaceutically acceptable salts thereof.
15

The individual enantiomers can be prepared by enantioselective synthesis from optically active precursors, or by resolving the racemic compound
20 which can be prepared as described above. Enantiomers of secondary amines of the formula I can also be prepared by preparing the racemate of the corresponding primary amine, resolving the latter into the individual enantiomers, and then converting the optically pure primary amine enantiomer into the required secondary amine by methods described in British Patent Specification
25 2098602.

Specific examples of compounds of formula I are:

(+)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N-methylamine;
30 (-)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N-methylamine;
(+)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
(-)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;

(+)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N,N-dimethylamine;
(-)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N,N-dimethylamine

or a pharmaceutically acceptable salt thereof.

5

The hydrochloride salts are preferred in each case, but the free bases and other pharmaceutically acceptable salts are also suitable.

10 The compound of formula I may be administered in any of the known pharmaceutical dosage forms. The amount of the compound to be administered will depend on a number of factors including the age of the patient, the severity of the condition and the past medical history of the patient and always lies within the sound discretion of the administering physician but it is generally envisaged that the dosage of the compound to be administered will be in the range 0.1 to 50
15 mg preferably 1 to 30 mg per day given in one or more doses.

Oral dosage forms are the preferred compositions for use in the present invention and these are the known pharmaceutical forms for such administration, for example tablets, capsules, granules, syrups and aqueous or oil suspensions.
20 The excipients used in the preparation of these compositions are the excipients known in the pharmacist's art. Tablets may be prepared from a mixture of the active compound with fillers, for example calcium phosphate; disintegrating agents, for example maize starch; lubricating agents, for example magnesium stearate; binders, for example microcrystalline cellulose or polyvinylpyrrolidone
25 and other optional ingredients known in the art to permit tabletting the mixture by known methods. The tablets may, if desired, be coated using known methods and excipients which may include enteric coating using for example hydroxypropylmethylcellulose phthalate. The tablets may be formulated in a manner known to those skilled in the art so as to give a sustained release of the
30 compounds of the present invention. Such tablets may, if desired, be provided with enteric coatings by known methods, for example by the use of cellulose acetate phthalate. Similarly, capsules, for example hard or soft gelatin capsules,

containing the active compound with or without added excipients, may be prepared by known methods and, if desired, provided with enteric coatings in a known manner. The contents of the capsule may be formulated using known methods so as to give sustained release of the active compound. The tablets and capsules may conveniently each contain 1 to 50 mg of the active compound, preferably 10 mg or 15 mg.

Other dosage forms for oral administration include, for example, aqueous suspensions containing the active compound in an aqueous medium in the presence of a non-toxic suspending agent such as sodium carboxymethylcellulose, and oily suspensions containing a compound of the present invention in a suitable vegetable oil, for example arachis oil. The active compound may be formulated into granules with or without additional excipients. The granules may be ingested directly by the patient or they may be added to a suitable liquid carrier (for example, water) before ingestion. The granules may contain disintegrants, eg an effervescent couple formed from an acid and a carbonate or bicarbonate salt to facilitate dispersion in the liquid medium.

The therapeutically active compounds of formula I may be formulated into a composition which the patient retains in his mouth so that the active compound is administered through the mucosa of the mouth.

Dosage forms suitable for rectal administration are the known pharmaceutical forms for such administration, for example, suppositories with cocoa butter or polyethylene glycol bases.

Dosage forms suitable for parenteral administration are the known pharmaceutical forms for such administration, for example sterile suspensions or sterile solutions in a suitable solvent.

Dosage forms for topical administration may comprise a matrix in which the pharmacologically active compounds of the present invention are dispersed

so that the compounds are held in contact with the skin in order to administer the compounds transdermally. A suitable transdermal composition may be prepared by mixing the pharmaceutically active compound with a topical vehicle, such as a mineral oil, petrolatum and/or a wax, e.g. paraffin wax or beeswax, together with
5 a potential transdermal accelerant such as dimethyl sulphoxide or propylene glycol. Alternatively the active compounds may be dispersed in a pharmaceutically acceptable cream, gel or ointment base. The amount of active compound contained in a topical formulation should be such that a therapeutically effective amount of the compound is delivered during the period
10 of time for which the topical formulation is intended to be on the skin.

The therapeutically active compound of formula I may be formulated into a composition which is dispersed as an aerosol into the patients oral or nasal cavity. Such aerosols may be administered from a pump pack or from a
15 pressurised pack containing a volatile propellant.

The therapeutically active compounds of formula I used in the method of the present invention may also be administered by continuous infusion either from an external source, for example by intravenous infusion or from a source of
20 the compound placed within the body. Internal sources include implanted reservoirs containing the compound to be infused which is continuously released for example by osmosis and implants which may be (a) liquid such as an oily suspension of the compound to be infused for example in the form of a very sparingly water-soluble derivative such as a dodecanoate salt or a lipophilic ester
25 or (b) solid in the form of an implanted support, for example of a synthetic resin or waxy material, for the compound to be infused. The support may be a single body containing all the compound or a series of several bodies each containing part of the compound to be delivered. The amount of active compound present in an internal source should be such that a therapeutically effective amount of the
30 compound is delivered over a long period of time.

In some formulations it may be beneficial to use the compounds of the present invention in the form of particles of very small size, for example as obtained by fluid energy milling.

- 5 In the compositions of the present invention the active compound may, if desired, be associated with other compatible pharmacologically active ingredients.

10 The invention further provides the use of compounds of formula I in the manufacture of a medicament for treating sexual dysfunction.

In another aspect, the invention further provides a pharmaceutical composition for treating sexual dysfunction, comprising a compound of formula I in conjunction with a pharmaceutically acceptable diluent or carrier.

- 15 The efficacy of compounds of formula I in treating sexual dysfunction is demonstrable through clinical trials in a relevant population set and the data presented below.

Monoamine reuptake inhibitors have been used to treat certain of the disorders described in the present invention. However, these compounds are known to suffer from a number of disadvantages. Firstly such compounds are 5 not effective in all patients. Secondly where the compounds are effective they may not provide a complete cure of the disorder. Thirdly, there are many undesirable side-effects known with this type of compound. Such side-effects include nausea, sexual dysfunction, light headedness, somnolence, sweating, tremor, dry mouth, asthenia, insomnia, diarrhoea, headache, vomiting, anxiety, 10 drowsiness, dizziness, fever, rash or allergic reactions, arthralgia, myalgia, convulsions, hypomania and mania.

Sibutramine (Formula I, $R_1 = CH_3$, $R_2 = CH_3$) has a pharmacological profile which is unique amongst monoamine reuptake inhibitors. Through its 15 pharmacologically active metabolites, (metabolite 1, $R_1 = H$, $R_2 = CH_3$ in Formula I and metabolite 2, $R_1 = H$, $R_2 = H$ in Formula I) sibutramine inhibits the reuptake of all three monoamines differentiating it from serotonin (5-HT)-selective reuptake inhibitors, e.g. fluoxetine, noradrenaline-selective reuptake inhibitors, e.g. desipramine, dopamine-selective reuptake inhibitors, e.g. bupropion, and 20 serotonin-noradrenaline reuptake inhibitors, e.g. venlafaxine (Table 1). It is this unique combination of pharmacological actions which renders sibutramine, and the other compounds of formula I, efficacious in the treatment of sexual dysfunction.

25 The assays below are performed in a similar manner to those described in WO98/41528.

TABLE 1

Comparison of the *in vitro* monoamine reuptake inhibition profiles of Examples 1
 5 and 2, and various reference monoamine reuptake inhibitors in rat brain tissue

	Ki (nM)		
	[³ H]Noradrenaline	[³ H]5-HT	[³ H]Dopamine
Example 1	3	18	24
Example 2	5	26	31
Bupropion	2590	18312	409
Desipramine	2	200	4853
Fluoxetine	320	11	2025
Venlafaxine	196	26	2594

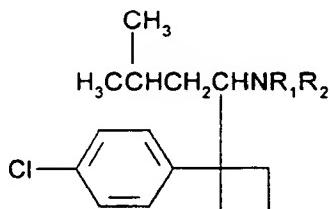
The results are the means of ≥ 3 separate determinations

Example 1 R₁ = H, R₂ = CH₃ in Formula I

Example 2 R₁ = H, R₂ = H in Formula I

Claims

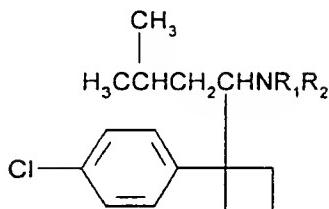
1. A method of treating sexual dysfunction comprising administering to a human in need thereof a therapeutically effective amount of a compound of
 5 formula I



including enantiomers and pharmaceutically acceptable salts thereof in which R₁ and R₂ are independently H or methyl, in conjunction with a pharmaceutically
 10 acceptable diluent or carrier.

2. A method as claimed in claim 1 wherein the compound of formula I is
N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride.
- 15 3. A method as claimed in claim 2 wherein the compound of formula I is
N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride in the form of its monohydrate.
- 20 4. A method as claimed in claim 1 in which the compound is selected from:
 (+)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N-methylamine;
 (-)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N-methylamine;
 (+)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
 (-)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
 25 (+)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N,N-dimethylamine;
 (-)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N,N-dimethylamine
 or a pharmaceutically acceptable salt thereof.

5. The use of a compound of formula I



5 including enantiomers and pharmaceutically acceptable salts thereof in which R₁ and R₂ are independently H or methyl, in the manufacture of a medicament for treating sexual dysfunction.

6. The use as claimed in claim 5 in which the compound of formula I is N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride.

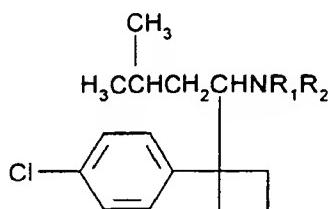
7. The use as claimed in claim 5 in which the compound of formula I is N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride monohydrate.

15 8. The use of a compound as claimed in claim 5 in which the compound is selected from:

- (+)-N-[1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl]-N-methylamine;
- 20 (-)-N-[1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl]-N-methylamine;
- (+)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
- (-)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
- (+)-N-[1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl]-N,N-dimethylamine;
- (-)-N-[1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl]-N,N-dimethylamine

25 or a pharmaceutically acceptable salt thereof.

9. A pharmaceutical composition for treating sexual dysfunction, comprising a therapeutically effective amount of a compound of formula I



5 including enantiomers and pharmaceutically acceptable salts thereof in which R₁ and R₂ are independently H or methyl, in conjunction with a pharmaceutically acceptable diluent or carrier.

10. A pharmaceutical composition as claimed in claim 9 in which the
10 compound of formula I is N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride.

11. A pharmaceutical composition as claimed in claim 9 in which the
15 compound of formula I is N,N-dimethyl-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine hydrochloride monohydrate.

12. A pharmaceutical composition as claimed in claim 9 in which the
compound of formula I is selected from:

20 (+)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N-methylamine;
(-)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N-methylamine;
(+)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
(-)-1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutylamine;
(+)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N,N-dimethylamine;
25 (-)-N-{1-[1-(4-chlorophenyl)cyclobutyl]-3-methylbutyl}-N,N-dimethylamine

or a pharmaceutically acceptable salt thereof.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US00/07114

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 31/135
US CL : 514/646

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/646

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
REG, structure search combined with erectile dysfunction or impotence or potent

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,459,164 A (VARGAS) 17 October 1995, see entire document.	5-12
X	US 4,939,175 A (UKAI et al.) 03 July 1990, see entire document.	5-12

Further documents are listed in the continuation of Box C.

See patent family annex.

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